	Ū	1		Document	ID	Issue Date	Pages
1			US	6123923	A	20000926	81
2			US	6120751	A	20000919	60
3			US	6090912	A	20000718	65
4			US	6090800	A	20000718	58
5			US	6028066	A	20000222	61
6			US	5840485	А	19981124	63
7			US	5232912	A	19930803	18
8			US	5086042	A	19920204	30
9			US	4472305 .	A	19840918	14

	Title	Current OR	Current XRef
1	Optoacoustic contrast agents and methods for their use	424/9.52	424/450 ; 424/9.1 ; 424/9.2 ; 424/9.3 ; 424/9.6 ; 514/410
2	Charged lipids and uses for the same	424/9.51	264/4 ; 264/4.1 ; 424/450 ; 424/502 ; 424/9.52 ; 428/402.2
3	Topologically segregated, encoded solid phase libraries comprising linkers having an enzymatically susceptible bond	530/300	435/212 ; 435/213 ; 436/518 ; 436/523 ; 436/528 ; 436/531 ; 530/304 ; 530/334 ; 530/402 ; 530/407
4	Lipid soluble steroid prodrugs	514/180	552/574
5	Prodrugs comprising fluorinated amphiphiles	514/180	514/169 ; 552/507
6	Topologically segregated, encoded solid phase libraries	435/6	435/7.1 ; 435/DIG.22 ; 435/DIG.34 ; 435/DIG.35 ; 435/DIG.38 ; 436/518 ; 530/300 ; 530/323 ; 536/23.1
7	Anticoagulant peptides	514/15	514/822 ; 530/328 ; 930/20 ; 930/21
8	Peptides with sulfate ester groups	514/16	514/15 ; 530/327 ; 530/328 ; 530/329
9	Hexapeptide amides	530/329	530/800 ; 930/10 ; 930/20 ; 930/21 ; 930/DIG.802 ; 930/DIG.803

	Retrieval Classif	Inventor	s	С	P	2	3	4	5
1		Unger, Evan C. , et al.	☒						
2		Unger, Evan C.	⊠						Ò
3		Lebl, Michal , et al.	×						
4		Unger, Evan C. , et al.	Ø						
5		Unger, Evan C.	⊠						
6		Lebl, Michal , et al.	⊠						
7		Krstenansky, John L.	×						
8		Rosamond, James D.	⊠						
9		Hansen, Philip E. , et al.	\boxtimes						

	U	1	Document	ID	Issue Date	Pages
10			US 4434096	А	19840228	5 '
11			US 4350627	Α	19820921	14
12			US 4028319	Α	19770607	11

	Title	Current OR	Current XRef
10	Substrates for the quantitative determination of proteolytic enzymes	530/331	
11	Biologically active peptides	530/302	514/809 ; 530/327 ; 530/328 ; 530/329 ; 930/10 ; 930/20 ; 930/21 ; 930/DIG.782 ; 930/DIG.803
12	2 AND 3-SUBSTITUTED ENKEPHALINS	530/302	514/809 ; 930/20 ; 930/21 ; 930/80 ; 930/DIG.741 ; 930/DIG.742

	Retrieval Classif	Inventor	S	С	P	2	3	4	5
10		Coleman, Patrick L. , et al.	☒						
11		de Castiglione, Roberto , et al.							
12	,	Jones, Jr., David A., et al.	⊠						

(FILE 'HOME' ENTERED AT 15:08:28 ON 06 FEB 2002)

```
FILE 'USPATFULL' ENTERED AT 15:08:50 ON 06 FEB 2002
            910 S (PHE OR TYR OR TRP) (P) (SAR OR AZE? OR NIP OR PIP OR PIPECOTIC
L1
            252 S (PHE OR TYR OR TRP) (2W) (SAR OR AZE? OR NIP OR PIP OR PIPECOTI
L2
L3
             48 S (PHE OR TYR OR TRP) (W) (PRO OR ALA OR GLY OR SER OR THR OR GLN
     FILE 'REGISTRY' ENTERED AT 15:14:50 ON 06 FEB 2002
              0 S [FYW]. [X]
L4
          24538 S [FYW].[X]/SQSP
L5
           1416 S L5 AND (SAR OR AZE OR NIP OR PIP)/NTE
L6
L7
            311 S L6 AND SQL<5
              0 S L6 AND SQL=3
L8
            303 S L6 AND SQL=4
1.9
           2927 S [STY]..[FYW].[X]/SQSP
L10
L11
            324 S L10 AND (SAR OR AZE OR NIP OR PIP)/NTE
            303 S L11 AND SQL=6
L12
     FILE 'CAPLUS' ENTERED AT 15:20:57 ON 06 FEB 2002
     FILE 'REGISTRY' ENTERED AT 15:21:25 ON 06 FEB 2002
           2119 S [STY][YGFMASILTVPKHQEZWRDNBC][YGFMASILTVPKHQEZWRDNBC][FYW].[X
L13
             24 S L13 AND (SAR OR AZE OR NIP OR PIP)/NTE
L14
              9 S L14 AND SQL=6
L15
     FILE 'CAPLUS' ENTERED AT 15:23:10 ON 06 FEB 2002
L16
             27 S L15
     FILE 'REGISTRY' ENTERED AT 15:27:17 ON 06 FEB 2002
            753 S [STY]..[FYW].[X]^/SQSP
L17
            303 S L17 AND (SAR OR AZE OR NIP OR PIP)/NTE
L18
     FILE 'CAPLUS' ENTERED AT 15:28:08 ON 06 FEB 2002
            427 S L18
L19
L20
              1 S 103:154282/DN
     FILE 'CAPLUS' ENTERED AT 15:32:40 ON 06 FEB 2002
     FILE 'USPATFULL' ENTERED AT 15:32:44 ON 06 FEB 2002
              0 S ALAPHATIC AMINO ACIDS
L21
              0 S ALAPHATIC AMINO
L22
L23
           1031 S ALIPHATIC AMINO ACID?
            347 S L23(P) TYR?
L24
L25
            248 S L23(P) (SER OR THR)
     FILE 'REGISTRY' ENTERED AT 15:35:59 ON 06 FEB 2002
            576 S [ST]..[FYW].[X]^/SQSP
L26
            299 S L26 AND (SAR OR AZE OR NIP OR PIP)/NTE
L27
            298 S L27 AND SQL=6
L28
=> s 128 and (cycl?)/cn
        423038 (CYCL?)/CN . . . .
L29
             0 L28 AND (CYCL?)/CN
=> s 128 not cycl?
       2823384 CYCL?
            81 L28 NOT CYCL?
L30
```

- u
- => s (phe or tyr or trp) (w) (pro or ala or gly or ser or thr or gln or asn or glu or asp or his or lys or arg or leu or ile or val or met or phe or try or trp or cys) (w) (sar or aze? or nip or pip or pipecotic or nipecotic)
 - 2 FILE BIOSIS
 - 1 FILE CANCERLIT
 - 14 FILES SEARCHED...
 - 16 FILE CAPLUS
 - 9 FILE DDFU
 - 1 FILE DGENE
 - 24 FILES SEARCHED...
 - 9 FILE DRUGU
 - 2 FILE EMBASE
 - 2 FILE ESBIOBASE
 - 34 FILES SEARCHED...
 - 2 FILE IFIPAT
 - 43 FILES SEARCHED...
 - 3 FILE MEDLINE
 - 2 FILE SCISEARCH
 - 54 FILES SEARCHED...
 - 1 FILE TOXLIT
 - 48 FILE USPATFULL
 - 58 FILES SEARCHED...
 - 4 FILE WPIDS
 - 4 FILE WPINDEX
 - 15 FILES HAVE ONE OR MORE ANSWERS, 61 FILES SEARCHED IN STNINDEX
- L35 QUE (PHE OR TYR OR TRP) (W) (PRO OR ALA OR GLY OR SER OR THR OR GLN OR ASN O R GLU OR ASP OR HIS OR LYS OR ARG OR LEU OR ILE OR VAL OR MET OR PHE O R TRY OR TRP OR CYS) (W) (SAR OR AZE? OR NIP OR PIP OR PIPECOTIC OR NIPE COTIC)

RL: SPN (Synthetic preparation); PREP (Preparation) (electrochem. redn. of the acyclic analog of pristinamycin Ia)

IT 182954-57-0P 182954-58-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (electrochem. redn. of the acyclic analog of pristinamycin Ia)

RN 182954-57-0 CAPLUS

CN Glycinamide, N-[(3-hydroxy-2-pyridinyl)carbonyl]-L-threonyl-(2R)-2-aminobutanoyl-L-prolyl-4-(dimethylamino)-N-methyl-L-phenylalanyl-(2S)-4-oxo-2-piperidinecarbonyl-2-phenyl-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 182954-58-1 CAPLUS

CN Glycinamide, N-[(3-methoxy-2-pyridinyl)carbonyl]-L-threonyl-(2R)-2-aminobutanoyl-L-prolyl-4-(dimethylamino)-N-methyl-L-phenylalanyl-(2S)-4-oxo-2-piperidinecarbonyl-2-phenyl-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

IT 182954-60-5P 182954-61-6P

RL: SPN (Synthetic preparation); PREP (Preparation) (electrochem. redn. of the acyclic analog of pristinamycin Ia) 182954-60-5 CAPLUS

CN Glycinamide, N-[(3-hydroxy-2-pyridinyl)methyl]-L-threonyl-(2R)-2-aminobutanoyl-L-prolyl-4-(dimethylamino)-N-methyl-L-phenylalanyl-(2S)-4-oxo-2-piperidinecarbonyl-2-phenyl-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN

RN 182954-61-6 CAPLUS

CN Glycinamide, N-[(3-methoxy-2-pyridinyl)methyl]-L-threonyl-(2R)-2-aminobutanoyl-L-prolyl-4-(dimethylamino)-N-methyl-L-phenylalanyl-(2S)-4-oxo-2-piperidinecarbonyl-2-phenyl-, (2S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L31 ANSWER 19 OF 29 CAPLUS COPYRIGHT 2002 ACS

AN 1996:641865 CAPLUS

DN 125:301589

TI Incidence of the peptidic lactone opening on the electrochemical reduction of pristinamycin IA

AU Largeron, Martine; Auzeil, Nicolas; Dakova, Bouria; Bacque, Eric; Paris, Jean-Marc; Fleury, Maurice-Bernard

CS Lab. Chimie Analytique Electrochimie, CNRS, Paris, 75270, Fr.

SO Tetrahedron Lett. (1996), 37(42), 7499-7502 CODEN: TELEAY; ISSN: 0040-4039

DT Journal

LA English

CC 34-3 (Amino Acids, Peptides, and Proteins) Section cross-reference(s): 72

OS CASREACT 125:301589

GΙ

AB Comparison of the cathodic behavior of pristinamycin IA with an open ring deriv. corroborates the role of the steric crowding exerted by the peptidic lactone. For example, ammonolysis of pristinamycin Ia produced I (R = H, CH3; X = CO) and preparative scale electrolysis of I (X = CO) produced I (X = CH2) in 30% yield.

Ι

ST electrochem redn pristinamycin acyclic analog; pristinamycin lactone ring opening ammonolysis

IT Reduction, electrochemical

(electrochem. redn. of the acyclic analog of pristinamycin Ia)

IT Ring cleavage

(of peptidic lactone, by ammonolysis; electrochem. redn. of the acyclic analog of pristinamycin Ia)

IT 3131-03-1D, Pristinamycin Ia, acyclic analogs

RL: RCT (Reactant)

(electrochem. redn. of the acyclic analog of pristinamycin Ia)

IT 182954-57-0P 182954-58-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (electrochem. redn. of the acyclic analog of pristinamycin Ia)

IT 182954-59-2P 182954-60-5P 182954-61-6P

```
L20
    ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
     1985:554282 CAPLUS
ΑN
     103:154282
DN
     Synthesis and study of N- and C-oligopeptide derivatives of
TI
     2-D-Ala,5-des-Met-enkephalin amide
     Vlasov, G. P.; Gusel, W. A.; Kozhevnikova, N. Yu.; Illarionova, N. G.;
ΑU
     Ditkovskaya, I. B.; Dorosh, M. Yu.; Krasnikova, E. N.
     Inst. Macromol. Comp., Leningrad, USSR
CS
     Pept., Proc. Eur. Pept. Symp., 18th (1984), 329-32. Editor(s):
SO
     Ragnarsson, Ulf. Publisher: Almqvist & Wiksell, Stockholm, Swed.
     CODEN: 53PWAN
DΤ
     Conference
     English
LΑ
     2-2 (Mammalian Hormones)
CC
     In a structure-activity study of 26 enkephalin analogs,
AΒ
     Tyr-D-Ala-Gly-Phe-D-Ala-NH2 [98537-64-5], Tyr-D-Ala-Gly-Phe-(D-Ala)2-NH2
     [98537-65-6], Tyr-D-Ala-Gly-Phe-(D-Ala)3-NH2 [98537-66-7], and
     Tyr-D-Ala-Gly-Phe-(D-Ala)4-NH2 [98537-67-8] had higher relative analgesic
     activity after intracerebroventricular administration of 5 mg into rats
     than did Tyr-D-Ala-Gly-Phe-NH2 (I) [66649-46-5]; whereas, the analgesic
     activity of 200 mg of (D-Ala)2-Tyr-D-Ala-Gly-Phe-NH2 [98537-59-8],
     (D-Ala)4-Tyr-D-Ala-Gly-Phe-NH2 [98537-60-1], or (D-Ala)6-Tyr-D-Ala-Gly-
     Phe-NH2 [98537-61-2] was only 25, 10, and 20% of that of I, resp. The
     effects of the enkephalins on body temp. is also given. Thus,
     modification of the N-terminus of I yields compds. with analgesic activity
     which is usually much lower than that of analogs modified at the
     C-terminus.
     enkephalin structure activity; analgesic enkephalin structure activity;
ST
     body temp enkephalin structure activity
IT
     Enkephalins
     RL: BIOL (Biological study)
        (analgesic activity and body temp. response to, structure in relation
        to)
ΙΤ
     Analgesics
        (enkephalins as, structure in relation to)
IT
     Body temperature
        (enkephalins effect on, structure in relation to)
     Molecular structure-biological activity relationship
IΤ
        (analgesic, of enkephalin analogs)
     Molecular structure-biological activity relationship
IT
        (body temp.-affecting, of enkephalin analogs)
                                            98537-59-8
                                                          98537-60-1
                  61090-95-7
                               66649-46-5
     58569-55-4
TΤ
                               98537-63-4
                                            98537-64-5
                                                          98537-65-6
     98537-61-2
                  98537-62-3
                  98537-67-8
                               98537-68-9
                                            98537-69-0
                                                          98537-70-3
     98537-66-7
                               98537-73-6
                                            98537-74-7
                                                          98537-75-8
                  98537-72-5
     98537-71-4
                               98537-78-1
                                            98537-79-2 . 98537-80-5
     98537-76-9
                  98537-77-0
     RL: BIOL (Biological study)
        (analgesic activity and body temp. response to, structure in relation
```

to)

L16 ANSWER 18 OF 27 CAPLUS COPYRIGHT 2002 ACS

AN 1985:554282 CAPLUS

DN 103:154282

TI Synthesis and study of N- and C-oligopeptide derivatives of 2-D-Ala,5-des-Met-enkephalin amide

AU Vlasov, G. P.; Gusel, W. A.; Kozhevnikova, N. Yu.; Illarionova, N. G.; Ditkovskaya, I. B.; Dorosh, M. Yu.; Krasnikova, E. N.

CS Inst. Macromol. Comp., Leningrad, USSR

Pept., Proc. Eur. Pept. Symp., 18th (1984), 329-32. Editor(s): Ragnarsson, Ulf. Publisher: Almqvist & Wiksell, Stockholm, Swed. CODEN: 53PWAN

DT Conference

LA English

IT 98537-74-7

RL: BIOL (Biological study)
(analgesic activity and body temp. response to, structure in relation to)

RN 98537-74-7 CAPLUS

CN Glycinamide, L-tyrosyl-D-alanylglycyl-L-phenylalanyl-N-methylglycyl-N2-methyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

```
L16 ANSWER 15 OF 27 CAPLUS COPYRIGHT 2002 ACS
```

AN 1987:423685 CAPLUS

DN 107:23685

TI Synthesis of enkephalin analogs on the polymer carrier Sephadex LH-20

AU Vlasov, G. P.; Gusel, V. A.; Kozhevnikova, N. Yu.; Ditkovskaya, I. B.; Dorosh, M. Yu.; Krasnikova, E. N.; Moskvicheva, Yu. B.; Kulyba, O. P.; Titov, A. P.; Kachurin, G. G.

CS Inst. Vysokomol. Soedin., Leningrad, USSR

SO Zh. Obshch. Khim. (1986), 56(7), 1635-41 CODEN: ZOKHA4; ISSN: 0044-460X

DT Journal

LA Russian

IT 98537-74-7P

RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation) (prepn. and analgesic activity of)

RN 98537-74-7 CAPLUS

CN Glycinamide, L-tyrosyl-D-alanylglycyl-L-phenylalanyl-N-methylglycyl-N2-methyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

IT 108787-67-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. and deprotection of)

RN 108787-67-3 CAPLUS

CN Glycinamide, N,O-bis[(1,1-dimethylethoxy)carbonyl]-L-tyrosyl-D-alanylglycyl-L-phenylalanyl-N-methylglycyl-N2-methyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B

IT 108787-66-2DP, Sephadex-bound

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and resin cleavage of, by aminolysis)

RN 108787-66-2 CAPLUS

CN Glycine, N-[N-[N-[N-[N-[N-[N,O-bis[(1,1-dimethylethoxy)carbonyl]-L-tyrosyl]-D-alanyl]glycyl]-L-phenylalanyl]-N-methylglycyl]-N-methyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-B